



SUNITINIB

INSTRUCTIONS FOR THE PHARMACIST

Prescription

- All orders should be written on a **pre-printed order**; if not, compare prescription to standard regimens in the Systemic Therapy Manual to confirm the dosing and instructions
 - The order must be signed by BOTH the prescriber (at the bottom) AND at least one other oncology health professional (nurse or hospital pharmacist) who has verified the order
- The prescription may **not** be refilled (unless specifically ordered by the oncologist) and it may **not** be filled as a continuing care prescription
 - If the prescriber has written for refills, contact the oncology team and do **not** dispense until they call back to authorize the refill; blood work must be checked for each cycle.
- **Always check for drug-drug interactions, especially before the first cycle. There is a strong potential for Sunitinib to interact with other drugs, foods or natural health products**, so a thorough drug interaction check (including medications filled at different pharmacies) is recommended before dispensing the first dose of Enzalutamide and after each new drug is considered for concomitant use. Consult the **Drug Interactions Table**, in this Toolkit.

Handling and Dispensing

- When handling this drug, disposable gloves should be worn at all times by any woman of child-bearing potential. Counting trays and other equipment directly exposed to the drug should be cleaned with **soap and water**, followed by rinsing with copious amounts of water (wear gloves).
- Do not crush tablets in an open air environment and risk inhalation of powder.
- ALWAYS affix the auxiliary label to identify this medication as “Cancer Chemotherapy”- this is an important warning label for other health professionals who care for the patient.



Patient Counseling and Follow-up

- Counsel the patient, including the key messages listed below. Use the **Initial Assessment and Patient Counseling Visit- Pharmacist Guide ①** and the **Medication Info Sheet ②** for this drug. Be sure that you know the specific treatment schedule and that this is clearly communicated to the patient.
- Call the patient within the first week to identify any problems with adverse reactions or adherence.
 - When speaking with the patient and during call backs, ask the patient to identify any problems with medication adherence. Use the **First Follow-Up Call/Visit- Pharmacist_Guide ①**.
- Continuing follow up calls between clinic visits are necessary for ADR identification and prevention and for adherence management. Contact the oncology clinic nurse or hospital pharmacist to negotiate who will do follow-up calls between clinic visits. Tell the patient that you plan to call back to check on their progress. Consider the suggested call-back schedule (pg. 2), with specific questions for each contact.
 - When speaking with the patient and during call backs, ask the patient to identify any problems with medication adherence. Use the **Continuing Follow-Up Calls/Visits - Pharmacist_Guide ①**.
- If the patient reports any adverse effects, consider the management strategies suggested in the **Adverse Drug Reaction Management Guide ②**.
- ALWAYS document your findings in the patient profile of your pharmacy computer system
- ALWAYS contact the patient’s cancer care team of any findings and actions you have taken.
- ALWAYS **watch for any unusual or unexpected symptoms or problems** (such as an adverse reaction that appears too soon or too severe) and contact the cancer care team promptly if something seems wrong with the patient experience.



CLINICAL INDICATIONS

SUNITinib is clinically indicated for:

- Gastrointestinal Stromal Tumour (GIST) after failure with Imatinib
- Advanced Renal Cell Carcinoma
- Pancreatic Neuroendocrine Tumours

DRUG ADMINISTRATION

- SUNITinib is generally taken **once daily**, with or without food. Do not crush or chew.
- Patients should not eat grapefruits or drink grapefruit juice while taking SUNITinib.
- Keep out of reach of children.
- There are different treatment schedules: One common schedule is to take the drug daily for 4 weeks, then take a 2 week break; for other indications the drug is taken continuously without a break.
- If a dose is missed, do not take a double dose the next day to make up for it.

PATIENT COUNSELLING- INITIAL AND FOLLOW-UP CALLS

In addition to other printed materials, use the **Medication Info Sheet** ② from the Cancer Care Nova Scotia website, and consider the more detailed suggestions in this toolkit.

	Key Messages
Initial counselling- At time of dispensing	<ul style="list-style-type: none"> • How to take the medication properly (including treatment-free breaks) • Symptoms to watch for reduced thyroid function • Prevention measures for Hand-Foot Skin Reaction and skin rashes • Good oral hygiene • When to call back to the cancer care team for urgent care • Use the Initial Assessment and Patient Counseling Visit- Pharmacist Guide ① and the drug-specific Medication Info Sheet ②
First call-back – Within first week: 	<ul style="list-style-type: none"> • Identify any initial problems with understanding or adherence • Use the First Follow-Up Call/Visit- Pharmacist Guide ① and the Medication Info Sheet ② (if needed) • Reinforce initial key messages <ul style="list-style-type: none"> ○ How and when treatment is taken ○ Barriers to adherence- remembering to take medication; reluctance to take treatment; financial issues; nausea, vomiting or other adverse effects; trouble with packaging; felt better off medication; other concerns ○ Suggest strategies to ensure adherence; reminder that full dose is needed for cancer control- partial doses may be ineffective. • Management of diarrhea- ensure patient has some Loperamide at home • Management of mild HFSR- ensure patient uses appropriate moisturizer • Identify any early adverse effect symptoms; suggest management strategies
Second call-back – After 2-3 weeks: (telephone or return visit to Pharmacy)	<ul style="list-style-type: none"> • Identify any adverse effects (PROBE for evidence of HFSR; skin rashes; changes in colour of skin or hair; symptoms of hypothyroidism, thrombosis, hypertension, CHF- trouble breathing, swollen ankles, QT prolongation-irregular heartbeats; mouth sores; diarrhea) or problems with understanding or adherence • Reinforce oral hygiene measures (ensure patient is following proper measures) • Management of mild HFSR- ensure patient uses appropriate moisturizer • Use the Continuing Follow-Up Calls/Visits- Pharmacist Guide ① <ul style="list-style-type: none"> ○ If any adverse effects identified, contact oncologist or oncology nurse

	Key Messages
End of Treatment call-back or 4 weeks from start:	<ul style="list-style-type: none"> Remind patient to stop taking pills after four weeks- ask if there are any pills left over; if so, PROBE to determine any barriers to treatment adherence Identify any adverse effects, as above
Subsequent cycles- (at least one call during each cycle:	<ul style="list-style-type: none"> Negotiate with patient and cancer care team for ongoing needs for counseling and timely follow up calls (every 1-3 months) Use the Continuing Follow-Up Calls/Visits- Pharmacist Guide 1 Adherence assessment and support is an important issue for reinforcement at each visit and mid-cycle call-back as treatment continues

ADVERSE EFFECTS: PREVENTION AND MANAGEMENT SUGGESTIONS

If you identify any of the following, you should contact the oncologist and tell the patient to call the oncologist or go directly to the Emergency Department of the nearest hospital right away **↗**:

- Blood clots (severe pain, swelling, or redness in legs or severe chest pain with shortness of breath)
- Heart problems (shortness of breath, fatigue, swollen feet and ankles)

NOTE- Thyroid function: SUNItinib can reduce thyroid function (symptoms: fatigue, anorexia, constipation, cold intolerance, dry skin, swelling or fluid retention, weight gain) as early as 2 weeks after therapy begins. Contact the cancer care team to arrange thyroid function testing if the patient reports any symptoms

The following are the common adverse effects from SUNItinib.

<p>More common</p> <p><u>Blood disorders</u></p> <ul style="list-style-type: none"> Low white blood cell (neutropenia) and platelet (thrombocytopenia) counts ★ <p><u>Cardiovascular disorders</u></p> <ul style="list-style-type: none"> Hypertension ★ ◆ <p><u>Gastrointestinal disorders</u></p> <ul style="list-style-type: none"> Mouth pain or irritation, stomatitis ★, taste disturbances, upset stomach, nausea ★, vomiting ★, diarrhea ★, constipation ★, loss of appetite <p><u>General disorders</u></p> <ul style="list-style-type: none"> Fatigue ★, headache ★ <p><u>Musculoskeletal disorders</u></p> <ul style="list-style-type: none"> Musculoskeletal pain ★- arthralgia, back pain, extremity pain <p><u>Skin disorders</u></p> <ul style="list-style-type: none"> Hand-foot skin reaction ★, skin rash ★, skin discolouration, hair-colour change ★ 	<p>Less Common</p> <p><u>Bleeding disorders</u></p> <ul style="list-style-type: none"> Nose bleed Venous thromboembolism (rare) ↗ <p><u>Cardiovascular disorders</u></p> <ul style="list-style-type: none"> Decrease cardiac function (LVEF) ↗ Myocardial infarction (rare) ↗ QT prolongation (rare) ↗ Venous thromboembolism (rare) ↗ <p><u>Gastrointestinal disorders</u></p> <ul style="list-style-type: none"> Abdominal pain <p><u>General disorders</u></p> <ul style="list-style-type: none"> Dizziness ◆, weakness Infection ◆ <p><u>Metabolic disorders</u></p> <ul style="list-style-type: none"> Hypothyroidism ↗; elevated liver enzymes (AST/ALT) ◆; elevated pancreatic enzymes (amylase, lipase) ◆; elevated creatinine ◆; hypokalemia ◆; hypernatremia ◆ <p><u>Skin disorders</u></p> <ul style="list-style-type: none"> Dry skin ★
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★ For detailed recommendations on the management of these adverse drug reactions, see the **Adverse Drug Reaction Management Guide 2**

◆ For management of these symptoms, the patient should see his physician

↗ These symptoms require urgent attention- advise the patient to go to the Emergency Department or contact their doctor (see instructions above)

DRUG INTERACTIONS

Take a **thorough medication history** (call other pharmacies if necessary) and determine the potential for all other drugs to increase or decrease SUNItinib plasma concentration.

- Drug interactions are often missed by community pharmacy computer systems
- **REPORT any potential interaction** to the prescribing oncologist- either the SUNItinib or the interaction drug may need to be dose altered or discontinued.

LIST OF IMPORTANT DRUG-DRUG INTERACTIONS WITH SUNITINIB- *This is not a complete list*

SUNItinib is metabolized primarily in the CYP3A4 pathway in the liver. Significant interactions are possible with other drugs that affect the same metabolic pathway.

- Anticoagulants (Anisindione, Ardeparin, Dalteparin, Dicoumarol, Enoxaparin, Heparin, Tinzaparin, Warfarin) - may increase risk of bleeding from Sunitinib
- Bisphosphonate agents (Alendronate, Etidronate, Ibandronate, Pamidronate, Risedronate, Zoledronic Acid)- may increase risk of osteonecrosis of the jaw from bisphosphonates
- **CYP 3A4 inducer** medications (e.g., dexamethasone, phenytoin, carbamazepine, rifampicin, phenobarbital or St. John’s Wort): consider dose increase of SUNItinib if co-administered with a strong CYP3A4 inducer (may decrease SUNItinib plasma concentrations)
- **CYP 3A4 inhibitor** medications (e.g., ketoconazole, itraconazole, erythromycin, clarithromycin): consider dose reduction of SUNItinib if co-administered with a strong CYP3A4 inhibitor (may increase SUNItinib plasma concentrations)
- Dabigatran- Increased levels of Dabigatrin in the blood
- Denosumab- Increased risk of serious infections
- Echinacea- Reduced SUNItinib levels
- Grapefruit or grapefruit juice- Increased SUNItinib blood levels
- Hepatotoxic drugs (Black Cohosh, Clofarabine, Interferon beta-1a, Interferon beta-1b, **Leflunomide**, Methotrexate, Naltrexone, **Teriflunomide**) - Increased risk of hepatotoxicity
- "Live" vaccinations
- PR prolongation- medications that cause a change in the heart rhythm
- QT prolongation- medications that cause a change in the heart rhythm
- Silodosin- Increased silodosin blood levels
- Tacrolimus ointment- Increased risk of serious infections, lymphoma and skin cancers

It is strongly recommended that you check any concurrent medications for interactions with this oral chemotherapy agent. Try one of the following comprehensive programs for checking drug interactions.

<p>Online Programs for Drug Interaction Checking-Publicly available:</p> <ul style="list-style-type: none"> • http://www.drugs.com/drug_interactions.php • http://reference.medscape.com/drug-interactionchecker • http://www.healthline.com/druginteractions • http://cpref.goldstandard.com/inter.asp?r=8084 • http://umm.edu/health/medical/drug-interaction-tool • http://online.epocrates.com/ (free account required) 	<p>Other Interaction Checkers- Subscription required:</p> <ul style="list-style-type: none"> • Lexicomp • Micromedex • eCPS
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